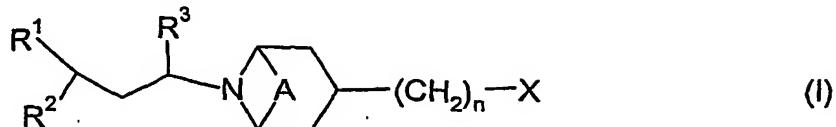


CLAIMS

1. A compound of formula (I):



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wherein:

A is absent or is $(CH_2)_2$;

R¹ is $C(O)NR^{10}R^{11}$, $C(O)_2R^{12}$, $NR^{13}C(O)R^{14}$, $NR^{15}C(O)NR^{16}R^{17}$, $NR^{18}C(O)_2R^{19}$, heterocyclyl (for example piperidine, piperazine, pyrrolidine or azetidine), aryl, cycloalkyl or heteroaryl;

10 R¹⁰, R¹³, R¹⁵, R¹⁶ and R¹⁸ are hydrogen or C₁₋₆ alkyl;

R¹¹, R¹², R¹⁴, R¹⁷ and R¹⁹ are C₁₋₈ alkyl (optionally substituted by halo, hydroxy, C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, C₃₋₆ cycloalkyl (optionally substituted by halo), C₅₋₆ cycloalkenyl, S(C₁₋₄ alkyl), S(O)(C₁₋₄ alkyl), S(O)₂(C₁₋₄ alkyl), heteroaryl, aryl, heteroaryloxy or aryloxy), aryl, heteroaryl, C₃₋₇ cycloalkyl (optionally substituted by halo or C₁₋₄ alkyl), C₄₋₇ cycloalkyl fused to a phenyl ring, C₅₋₇ cycloalkenyl, or, heterocyclyl (itself optionally substituted by oxo, C(O)(C₁₋₆ alkyl), S(O)_k(C₁₋₆ alkyl), halo or C₁₋₄ alkyl); or R¹¹, R¹², R¹⁴ and R¹⁷ can also be hydrogen;

15 or R¹⁰ and R¹¹, and/or R¹⁶ and R¹⁷ may join to form a 4-, 5- or 6-membered ring which optionally includes a nitrogen, oxygen or sulphur atom, said ring being optionally substituted by C₁₋₆ alkyl, S(O)_l(C₁₋₆ alkyl) or C(O)(C₁₋₆ alkyl);

20 R² is phenyl, heteroaryl or C₃₋₇ cycloalkyl;

R³ is H or C₁₋₄ alkyl;

X is $S(O)_2NR^4R^5$ or $NR^6S(O)_2R^7$;

25 R⁷ is aryl, heteroaryl, C₁₋₆ alkyl, C₃₋₇ cycloalkyl, heterocyclyl or NR^8R^9 wherein NR^8R^9 can be cyclized to form a 4-, 5- or 6-membered ring which optionally includes a nitrogen, oxygen or sulphur atom, said ring being optionally substituted by C₁₋₆ alkyl, S(O)_p(C₁₋₆ alkyl) or C(O)(C₁₋₆ alkyl);

R⁴ and R⁸ are aryl, heteroaryl, C₁₋₆ alkyl (optionally substituted by hydroxy or C₁₋₆ alkoxy), C₃₋₇ cycloalkyl or heterocyclyl;

30 R⁵, R⁶ and R⁹ are, independently, hydrogen or C₁₋₆ alkyl;

n is 1, 2 or 3;

aryl, phenyl and heteroaryl moieties are independently optionally substituted by one or more of halo, cyano, nitro, hydroxy, OC(O)NR²⁰R²¹, NR²²R²³, NR²⁴C(O)R²⁵,

NR²⁶C(O)NR²⁷R²⁸, S(O)₂NR²⁹R³⁰, NR³¹S(O)₂R³², C(O)NR³³R³⁴, CO₂R³⁶,

NR³⁷CO₂R³⁸, S(O)_qR³⁹, OS(O)₂R⁴⁹, C₁₋₆ alkyl (optionally mono-substituted by

S(O)₂R⁵⁰ or C(O)NR⁵¹R⁵²), C₂₋₆ alkenyl, C₂₋₆ alkynyl, C₃₋₁₀ cycloalkyl, C₁₋₆ haloalkyl,

C₁₋₆ alkoxy(C₁₋₆)alkyl, C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, phenyl, phenyl(C₁₋₄)alkyl,

phenoxy, phenylthio, phenylS(O), phenylS(O)₂, phenyl(C₁₋₄)alkoxy, heteroaryl,

heteroaryl(C₁₋₄)alkyl, heteroaryloxy or heteroaryl(C₁₋₄)alkoxy; wherein any of the

immediately foregoing phenyl and heteroaryl moieties are optionally substituted with

halo, hydroxy, nitro, S(C₁₋₄ alkyl), S(O)(C₁₋₄ alkyl), S(O)₂(C₁₋₄ alkyl), S(O)₂NH₂,

S(O)₂NH(C₁₋₄ alkyl), S(O)₂N(C₁₋₄ alkyl)₂, cyano, C₁₋₄ alkyl, C₁₋₄ alkoxy, C(O)NH₂,

C(O)NH(C₁₋₄ alkyl), C(O)N(C₁₋₄ alkyl)₂, CO₂H, CO₂(C₁₋₄ alkyl), NHC(O)(C₁₋₄ alkyl),

NHS(O)₂(C₁₋₄ alkyl), CF₃ or OCF₃;

unless otherwise stated heterocyclyl is optionally substituted by C₁₋₆ alkyl [optionally substituted by phenyl {which itself optionally substituted by halo, C₁₋₄ alkyl, C₁₋₄

alkoxy, cyano, nitro, CF₃, OCF₃, (C₁₋₄ alkyl)C(O)NH, S(O)₂NH₂, C₁₋₄ alkylthio,

S(O)(C₁₋₄ alkyl) or S(O)₂(C₁₋₄ alkyl)} or heteroaryl {which itself optionally substituted

by halo, C₁₋₄ alkyl, C₁₋₄ alkoxy, cyano, nitro, CF₃, (C₁₋₄ alkyl)C(O)NH, S(O)₂NH₂, C₁₋₄

alkylthio, S(O)(C₁₋₄ alkyl) or S(O)₂(C₁₋₄ alkyl)}], phenyl {optionally substituted by

halo, C₁₋₄ alkyl, C₁₋₄ alkoxy, cyano, nitro, CF₃, OCF₃, (C₁₋₄ alkyl)C(O)NH, S(O)₂NH₂,

C₁₋₄ alkylthio, S(O)(C₁₋₄ alkyl) or S(O)₂(C₁₋₄ alkyl)}, heteroaryl {optionally substituted

by halo, C₁₋₄ alkyl, C₁₋₄ alkoxy, cyano, nitro, CF₃, (C₁₋₄ alkyl)C(O)NH, S(O)₂NH₂, C₁₋₄

alkylthio, S(O)(C₁₋₄ alkyl) or S(O)₂(C₁₋₄ alkyl)}, S(O)₂NR⁴⁰R⁴¹, C(O)R⁴², C(O)₂(C₁₋₆

alkyl) (such as tert-butoxycarbonyl), C(O)₂(phenyl(C₁₋₂ alkyl)) (such as

benzyloxycarbonyl), C(O)NHR⁴³, S(O)₂R⁴⁴, NHS(O)₂NHR⁴⁵, NHC(O)R⁴⁶,

NHC(O)NHR⁴⁷ or NHS(O)₂R⁴⁸, provided none of these last four substituents is linked

to a ring nitrogen;

k, l, p and q are, independently, 0, 1 or 2;

R²⁰, R²², R²⁴, R²⁶, R²⁷, R²⁹, R³¹, R³³, R³⁷, R⁴⁰ and R⁵¹ are, independently, hydrogen or

C₁₋₆ alkyl;

R²¹, R²³, R²⁵, R²⁸, R³⁰, R³², R³⁴, R³⁶, R³⁸, R³⁹, R⁴¹, R⁴², R⁴³, R⁴⁴, R⁴⁵, R⁴⁶, R⁴⁷, R⁴⁸,

R⁴⁹, R⁵⁰ and R⁵² are, independently, C₁₋₆ alkyl (optionally substituted by halo, hydroxy,

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C₁₋₆ alkoxy, C₁₋₆ haloalkoxy, C₃₋₆ cycloalkyl, C₅₋₆ cycloalkenyl, S(C₁₋₄ alkyl), S(O)(C₁₋₄ alkyl), S(O)₂(C₁₋₄ alkyl), heteroaryl, phenyl, heteroaryloxy or phenoxy, C₃₋₇ cycloalkyl, phenyl or heteroaryl; wherein any of the immediately foregoing phenyl and heteroaryl moieties are optionally substituted with halo, hydroxy, nitro, S(C₁₋₄ alkyl), S(O)(C₁₋₄ alkyl), S(O)₂(C₁₋₄ alkyl), S(O)₂NH₂, S(O)₂NH(C₁₋₄ alkyl), S(O)₂N(C₁₋₄ alkyl)₂, cyano, C₁₋₄ alkyl, C₁₋₄ alkoxy, C(O)NH₂, C(O)NH(C₁₋₄ alkyl), C(O)N(C₁₋₄ alkyl)₂, CO₂H, CO₂(C₁₋₄ alkyl), NHC(O)(C₁₋₄ alkyl), NHS(O)₂(C₁₋₄ alkyl), C(O)(C₁₋₄ alkyl), CF₃ or OCF₃;

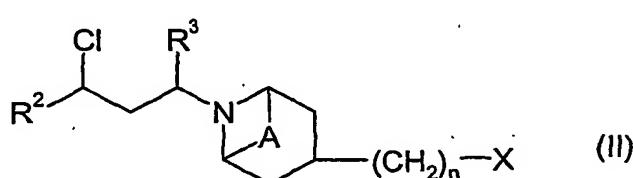
5 R²¹, R²³, R²⁵, R²⁸, R³⁰, R³⁴, R³⁵, R³⁶, R⁴¹, R⁴², R⁴³, R⁴⁵, R⁴⁶, R⁴⁷ and R⁵² may additionally be hydrogen;

10 or a pharmaceutically acceptable salt thereof or a solvate thereof.

2. A compound as claimed in claim 1 wherein A is absent.
- 15 3. A compound as claimed in claim 1 or 2 wherein n is 1 or 2.
4. A compound as claimed in claim 1, 2 or 3 wherein R³ is hydrogen.
5. A compound as claimed in claim 1, 2, 3 or 4 wherein R¹ is NR¹³C(O)R¹⁴; wherein R¹³ and R¹⁴ are as defined in claim 1.
- 20 6. A compound as claimed in claim 1, 2, 3 or 4 wherein R¹ is optionally substituted aryl or optionally substituted heteroaryl, wherein the optional substituents are as recited in claim 1.
- 25 7. A compound as claimed in claim 1, 2, 3 or 4 wherein R¹ is optionally substituted heterocyclyl.
8. A compound as claimed in any one of the preceding claims wherein R² is phenyl optionally substituted by halo or CF₃.
- 30 9. A compound as claimed in any one of the preceding claims wherein X is NR⁶S(O)₂R⁷; wherein R⁶ and R⁷ are as defined in claim 1.

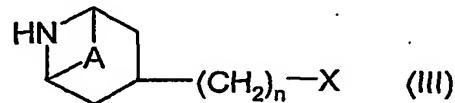
10. A compound as claimed in any one of the preceding claims wherein X is $S(O)_2NR^4R^5$;
 wherein R^4 and R^5 are as defined in claim 1.

5 11. A process for preparing a compound as claimed in claim 1, the process comprising:
 a. when R^1 is an N-linked optionally substituted heterocycle, reacting a compound
 of formula (II):

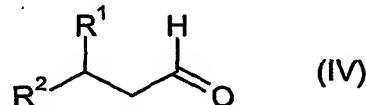


10 wherein R^2 , R^3 , n, A and X are as defined in claim 1, with a compound R^1H
 (wherein the H is on a heterocycle ring nitrogen atom) wherein R^1 is as defined
 above, in the presence of a suitable base, in a suitable solvent and optionally in
 the presence of sodium iodide;

b. when R^3 is hydrogen, coupling a compound of formula (III):

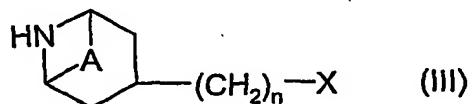


15 wherein n, A and X are as defined in claim 1, with a compound of formula (IV):



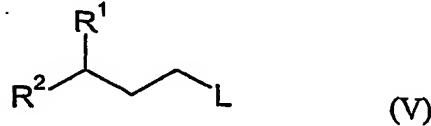
wherein R^1 and R^2 are as defined in claim 1, in the presence of $NaBH(OAc)_3$ in a
 suitable solvent at room temperature;

c. when R^3 is hydrogen, coupling a compound of formula (III):



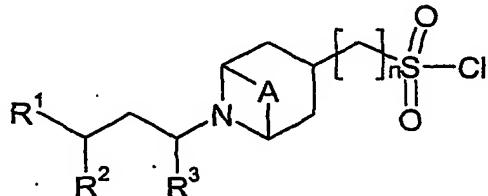
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wherein n, A and X are as defined in claim 1, with a compound of formula (V):



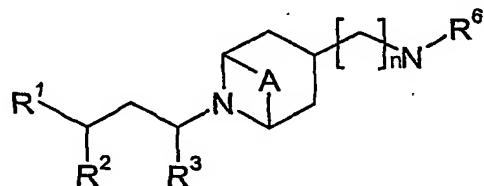
wherein R¹ and R² are as defined in claim 1 and L is a leaving group; in the presence of a base, in a suitable solvent at a temperature from 60°C up to the boiling point of the solvent;

d. when X is S(O)₂NR⁴R⁵, reacting a compound:



wherein R¹, R², R³, A and n are as defined in claim 1, with NHR⁴R⁵, wherein R⁴ and R⁵ are as defined in claim 1, in the presence of a suitable base and in the presence of a suitable solvent; or,

e. when X is NR⁶S(O)₂NR⁷, reacting a compound:



wherein R¹, R², R³, A and n are as defined in claim 1, with R⁷S(O)₂Cl, in the presence of a suitable base and in the presence of a suitable solvent.

12. A pharmaceutical composition which comprises a compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof, and a pharmaceutically acceptable adjuvant, diluent or carrier.

13. A compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof, for use as a medicament.

14. A compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof, in the manufacture of a medicament for use in therapy.

15. A method of treating a CCR5 mediated disease state comprising administering to a patient in need of such treatment an effective amount of a compound as claimed in claim 1, or a pharmaceutically acceptable salt thereof or solvate thereof.